THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants:

Jordan J.N. Tang and Arun K. Ghosh

TECH CENTER 1600/2900

Serial No.:

09/506,988

Art Unit:

1625

Filed:

February 18, 2000

Examiner:

Seaman, D.

For:

PROTEASE INHIBITORS THAT OVERCOME DRUG RESISTANCE

Assistant Commissioner for Patents Washington, D.C. 20231

SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT

Sir:

Pursuant to the duty of disclosure under 37 C.F.R. §1.56 and 37 C.F.R. §1.97, Applicants submit a Supplemental Information Disclosure Statement, including two (2) pages of Form PTO-1449, copies of the documents cited therein, and a copy of the International Search Report. mailed 13 November 2000, in the corresponding PCT application PCT/US00/04215.

Enclosed is a check for \$240.00 representing the fee required under 37 C.F.R. §1.17(p) for an Information Disclosure Statement filed after a first office action on the merits under 37 C.F.R. §1.97(c). It is believed that no additional fees are required with this submission. However, should a fee be required, the Commissioner is hereby authorized to charge any fees to

Deposit Account No. 01-2507.

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Hoechst Aktiengesellschaft

Country EP

WO 92/03472 A1

03-05-1992

The Upjohn Company

PCT

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the art referred to herein, either alone or in combination.

SUPPLEMENTAL INFORMATION

DISCLOSURE STATEMENT

Remarks

This statement should not be interpreted as a representation that an exhaustive search has been conducted or that no better art exists. Moreover, Applicants invite the Examiner to make an independent evaluation of the cited art to determine its relevance to the subject matter of the present application. Applicants are of the opinion that their claims patentably distinguish over

Respectfully submitted,

Patrea L. Pabst

Reg. No. 31,284

Dated: February 22, 2001

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SUPPLEMENTAL INFORMATION

DISCLOSURE STATEMENT

Publications

BAKER & CONDON, "Dipeptide isosteres. 1. Synthesis of dihydroxyethylene dipeptide isosteres via diastereoselective additions of alkyllithium reagents to *N*,*N*-dimethylhydrazones. Preparations of renin and HIV-1 protease inhibitors transition-state mimics," *J Org Chem* 58:3277-3284 (1993).

BAKER, et al., "Nonpeptide renin inhibitors employing a novel 3-Aza (or oxa)-2,4-dialkyl glutaric acid moiety as a P_2/P_3 amide bond replacement," *J Med Chem* 35:1722-1734 (1992).

BENNETT, et al., "The synthesis of novel HIV-protease inhibitors via silica gel assisted addition of amines to epoxides," SYNLETT 9:703-704 (1993).

DREYER, et al., "Inhibition of human immunodeficiency virus 1 protease in vitro: rational design of design of substrate analogue inhibitors," *Proc Natl Acad Sci. USA* 86:9752-9756 (1989).

MARINIER, et al., "HIV-1 protease inhibitors: ketomethylene isosteres with unusually high affinity compared with hydroxyethylene isostere analogs," *Bioorganic & Medicinal Chemistry* 2(9):919-925 (1994).

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DISCLOSURE STATEMENT

Certificate of Mailing under 37 CFR §1.8(a)

I hereby certify that this Supplemental Information Disclosure Statement, along with any paper referred to as being attached or enclosed, is being deposited with the United States Postal Service on the date shown below with sufficient postage as first-class mail in an envelope addressed to the Assistant Commissioner for Patents, Washington, D.C. 20231.

Date:

February 22, 2001